CLAIMS

What is claimed is:

1. A compound of the formula:

5 wherein:

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 R^1 is selected from C_1 - C_{10} alkyl, C_2 - C_{10} alkenyl, or C_2 - C_{10} alkynyl optionally substituted by OH, alkoxy, phenoxy, thio C_1 - C_{10} alkyl, or NR^4R^5 ; (CH₂)n-Ar, wherein the (CH₂)_n alkyl chain is optionally substituted by OH, alkoxy, phenoxy, thio C_1 - C_{10} alkyl, or NR^4R^5 ; COR⁴, wherein R^4 is alkyl optionally substituted by OH, alkoxy, phenoxy, thio C_1 - C_{10} alkyl, or NR^4R^5 ; C_3 - C_{10} cycloalkyl optionally substituted by OH, alkoxy, phenoxy, NR^4R^5 , SO₂ NR^4R^5 , or SO₃ R^4 ; (CH₂)_nheterocyclyl; or alkyl optionally substituted by COR⁴, CO_2R^4 or CONR⁴ R^5 ;

R⁴ is H or C₁-C₆ alkyl;

 R^5 is C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, $(CH_2)_nAr$, C_3 - C_{10} cycloalkyl, heterocyclyl or heteroaryl

n is 0 to 3;

 R^3 is $(CH_2)_nAr$;

Ar is phenyl optionally substituted by halo or alkyl optionally substituted by OH, alkoxy, phenoxy, thio C_1 - C_{10} alkyl, or NR^4R^5 ;

 R^2 is hydrogen; C_1 - C_{10} alkyl substituted by halo, nitrile, OH, alkoxy, phenoxy, thio C_1 - C_{10} alkyl, NR^4R^5 or (CH_2) -heteroaryl; $(CH_2)_nAr$, wherein n is 0-3; -(CH₂)-heteroaryl; C_3 - C_{10} cycloalkyl optionally substituted by OH, alkoxy, phenoxy, NR^4R^5 , $SO_2NR^4R^5$, or SO_3R^4 ; (CH_2) -heterocyclyl; or COR^4 ;

 R^4 is H, C_1 - C_6 alkyl optionally substituted by halogen; NR^5R^6 ; cycloalkyl; or (CH₂)-Ar;

 R^5 and R^6 are independently C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, $(CH_2)_nAr$, C_3 - C_{10} cycloalkyl, heterocyclyl or heteroaryl; or a pharmaceutically acceptable salt form thereof.

- 2. A compound of Claim 1 wherein R^3 is $(CH_2)_nAr$ substituted by one or two halogens.
- 3. A compound of Claim 1 wherein R^2 is hydrogen; C_1 - C_{10} alkyl optionally substituted by halo, nitrile, OH, alkoxy, phenoxy, thio C_1 - C_{10} alkyl, NR^4R^5 or (CH_2) -heteroaryl.
- 4. A method for the preparation of a compound of Claim 1, said method comprising:
 - (a) treating a compound of the formula:

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$$\begin{bmatrix} \\ \\ \\ \\ \\ \\ \\ \end{bmatrix}$$

wherein L is a leaving group, with an amine of the formula R^1 -NH₂, wherein n, R^1 , R^2 and R^3 have the meanings provided in Claim 1.

- 5. A method for the preparation of a compound of Claim 1, said method comprising:
- 25 (a) treating a compound of the formula:

with an oxidizing agent followed by an amine of the formula R^1 -NH₂, wherein n, R^1 , R^2 and R^3 have the meanings provided in Claim 1.

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6. A pharmaceutical composition comprising a pharmaceutically effective amount of a compound of Claim 1 and a pharmaceutically acceptable carrier.

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